WHAT IS CLAIMED IS:

1. A compound selected from the group consisting of compounds of the formula

NHR R_{4} R_{2} C = N R_{4} R_{2} NR_{3} C = N R_{4} R_{2} NR_{3} NHR_{1} R_{4} R_{2} NHR_{1} R_{4} R_{2} NHR_{1} R_{4} R_{2}

wherein R is a lower alkyl group of 1 to 6 carbon atoms;

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 R_1 is hydrogen or a lower alkyl group of 1-6 carbon atoms, amino, or hydroxy, or together with R_2 represent a lower alkylene bridge of 2-4 carbon atoms;

 R_2 is hydrogen, or a lower alkyl group of 1-6 carbon atoms, a hydroxyethyl group or R_2 may be taken together with R_1 as noted above;

- R_3 is hydrogen, a lower alkyl group of 1-6 carbon
- 37 atoms or may be together with R₁ a lower alkylene bridge
- 38 of 2-4 carbon atoms;
- and R_4 is hydrogen, a lower alkyl group of 1-6 carbon
- 40 atoms or together with R₃ is a lower alkylene bridge of 2-
- 41 4 carbon atoms; and their pharmaceutically acceptable
- 42 salts.
 - 1 2. A compound according to Claim 1 wherein R is lower
 - 2 alkyl and R_1 , R_2 , R_3 and R_4 are each hydrogen, and their
 - 3 pharmaceutically acceptable salts.
 - 1 3. The compound according to Claim 1 which is 3-amino-
 - 2 5-propylaminomethyl-6-2',3'-dihydroxypropyl)-1,2,4-
 - 3 triazine or a pharmaceutically acceptable salt thereof.
 - 1 4. The compound according to Claim 1 which is 1-
 - 2 propylamino-2,3-diaminoguanidine-1,4-dideoxyglucosone
 - 3 dihydrazone or a pharmaceutically acceptable salt
 - 4 thereof.
 - 1 5. A test kit for the detect on of the glycosylation
 - 2 products of polypeptides,//comprising:
 - a. a predetermined amount of a labeled compound of
 - 4 formula I or II or the binging partner specific thereto;
 - 5 b. other reagents; and
 - 6 c. directions for use of said kit.
 - 1 6. A test kit to be used for the detection and/or
 - 2 determination of one of the components selected from the
 - 3 group consisting of glycosylation products of
 - 4 polypeptides, and the specific binding partners thereto,
 - 5 according to a predetermined protocol, comprising:
 - a. a labeled component which has been obtained by
 - 7 coupling a compound of formula I or II to a detectable
 - 8 label;

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one or more additional immunochemical reagents b. 9 of which at least one reagent is a ligand or an 10 immobilized ligand, which ligand is selected from the 11 group consisting of: 12 a ligand capable/of binding with the 13 14 labeled component (a); (ii) a ligand capable of binding with a binding 15 partner of the labeled component (a); 16 (iii) a ligand/capable of binding with at least 17 one of the component(s) to be determined; and 18 (iv) a ligard/capable of binding with at least 19 one of the binding parkners of at least one of the 20 component(s) to be determined; and 21 directions for the performance of a protocol 22 for the detection and/or determination of one or more 23 components of an immunochemical reaction between the 24 advanced glycosylation end product and a specific binding 25

An indicator composition for use in an assay procedure for the detection of advanced glycosylation endproducts in polypeptide samples, said composition 3 4 comprising a compound selected from the group consisting of compounds of the formula

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NHR NHR.

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24 C = N - NNHR C = N - NOH C = N - NOH

wherein R is a lower alkyl group of 1 to 6 carbon atoms;

 R_1 is hydrogen or a lower alkyl group of 1-6 carbon atoms, amino, or hydroxy, or together with R_2 represent a lower alkylene bridge of 2-4 carbon atoms;

 R_2 is hydrogen, or a lower alkyl group of 1-6 carbon 40 atoms, a hydroxyethyl group or R_2 may be taken together 41 with R_1 as noted above;

 R_3 is hydrogen, a lower alkyl group of 1-6 carbon atoms or may be together with R_1 a lower alkylene bridge of 2-4 carbon atoms;

and R_4 is hydrogen, a lower alkyl group of 1-6 carbon 46 atoms or together with R_3 is a lower alkylene bridge of 2-47 4 carbon atoms; and their pharmaceutically acceptable 48 salts.

An indicator composition according to Claim of for use in an assay procedure for the detection of advanced glycosylation endproducts in polypeptide samples, said pomposition comprising a compound which is a 3-amino-5-

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5 alkylaminomethyl-6-alkyl-1,2,4-triazine of the formula (I/a)

- 1 9. The indicator of Claim 7 having associated therewith
- 2 a detectable label.
- 1 10. The indicator of Claim 9 wherein the label is an
- 2 enzyme.
- 1 11. The indicator of Claim 10/wherein the label is
- 2 selected from peroxidase, β-g/lucuronidase, β-D-
- 3 glucosidase, B-D-galactosidase, urease, glucose oxidase
- 4 plus peroxidase, galactose oxidase plus peroxidase, and
- 5 acid phosphatase.
- 1 12. The indicator of Claim 9 wherein said label is a
- 2 radioactive element.
- 1 13. The indicator/of/Claim /12 wherein said radioactive
- 2 element is selected from the group consisting of 14C, 125I,
- $3^{131}I$, ^{35}S and ^{3}H .
- 1 14. The indicator of claim 9 wherein said label is a
- 2 chemical which fluoresces when exposed to ultraviolet
- 3 light.
- 1 15. The indicator of Claim 14 wherein said chemical is
- 2 selected from fluorescein, rhodamine, and auramine.
- 1 16. A method for the preparation of compounds of
- 2 formulae I and II comprising non-enzymatically reacting,
- 3 under physiological conditions, a 1-alkylamino-1,4-
- 4 dideoxyosone of the formula

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- 1 20. A method for measuring the amount of advanced
- 2 glycosylation endproducts in a protein sample comprising
- 3 measuring the presence and amount/of a compound of Claim
- 4 1.
- 1 21. A method of cross linking proteins by reacting said
- 2 proteins with a compound of Claim 1.
- 1 22. A method of quantitating proteins in a biological
- 2 sample by measuring the reactivity of the proteins with a
- 3 known amount of a compound of Claim 1.
- 1 23. A method of increasing the immunogenicity of an
- 2 antigen which comprises crosslinking said antigen with a
- 3 compound of Clayim 1.
- 1 24. A composition for promoting the sequestration and
- 2 removal from the body of an animal of target
- 3 macromolecules that have undergone advanced glycosylation
- 4 comprising a compound of Claim 1 capable of causing the
- 5 body to increase its activity of recognizing and removing
- 6 said macromolecules.
- 1 25. The composition of Claim 47 wherein said compound is

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- 2 bound to a carrier.
- 1 26. A method for the preparation of advanced
- 2 glycosylation endproducts which comprises the
- 3 nonenzymatic reaction of a compound of the formula

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10 wherein R is a lower alky group, under physiological

11 conditions.

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wherein R is a lower alkyl group; 9 with aminoguanidine or an analog of the formula 10 11 (IVb) R4HN-N-C-NHR1 12 13 NR₃ 14 15 wherein R is hydrogen or a lower alkyl group of 1-6 16 carbon atoms, amino, or hydroxy, or together with R2 17 represent a lower alkylene bridge of 2-4 carbon atoms; 18 R₂ is hydrogen, or a lower alkyl group of 1-6 carbon 19 atoms, a hydroxyethyl group or R_2 may be taken together 20 with R₁ as noted above; 21 R₁ is hydrogen, a lower alkyl group of 1-6 carbon 22 atoms or may be together with R₁ a lower alkylene bridge 23 of 2-4 carbon atoms; and 24 R₄ is hydrogen, a lower alkyl group of 1-6 carbon 25 atoms or together with R_3^{\prime} is a lower alkylene bridge of 2-26 27 4 carbon atoms; and their pharmaceutically acceptable salts. 28 A method for measuring the amount of aminoguanidine 1 or its analogs in /a/protein sample comprising measuring 2 the presence and amount of a compound of Claim 1. 3 A method of detecting an aminoguanidine allergy in 1 humans comprising testing the serum of the patient to 2 determine the présence of antibodies to a compound of 3 4 Claim 1. A method for removing advanced glycosylation 1 endproducts from the body by administering the anti-2 antibody or second binding partner to a compound of Claim 3 1 to form an/immune complex activating the animal's 4 cellular clearance system (macrophages) to remove said 5 immune complex and associated AGEs (advanced 6 glycosylation endproducts).